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- a) bringing the sample into contact with an *H. pylori* bacterial strain having an aflagellate phenotype resulting from a mutation in the *flbA* gene of said *H. pylori* bacterial strain such that either the *flbA* gene is no longer expressed in the *H. pylori* bacterial strain, or the expression of the *flbA* gene in the *H. pylori* bacterial strain does not enable the A and B flagellins or the sheath that contains them to be biosynthesized and, if this is the case, does not enable the *H. pylori* anchoring protein or the hook to be synthesized; and
- b) detecting an immunological reaction between the bacterial strain and antibodies directed against *H. pylori* and which are present in the sample.
- 44. The method as claimed in claim 43, wherein the aflagellate *H. pylori* strain also does not express the hook protein of *H. pylori*.
- 45. The method as claimed in claim 43, wherein the *H. pylori* bacterial strain is obtained from strain N6 having deposit Accession No. NCIMB 40512.
- 46. The method as claimed in claim 43, wherein the aflagellate strain of *H. pylori* is strain N6flbA⁻ having deposit Accession No. NCIMB 40747.
- 47. The method as claimed in claim 43, wherein the biological fluid is human serum, saliva, or urine.
- 48. The method as claimed in claim 43, wherein the immunological reaction is detected by Western blot or ELISA.
- 49. Method for the *in vitro* detection of an infection due to *H. pylori* in a sample of biological fluid from a patient, wherein the method comprises:

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- bringing the sample into contact with a <u>bacterial extract</u> from an *H. pylori* bacterial strain having an aflagellate phenotype resulting from a mutation in the *flbA* gene of said *H. pylori* bacterial strain such that either the *flbA* gene is no longer expressed in the *H. pylori* bacterial strain, or the expression of the *flbA* gene in the *H. pylori* bacterial strain does not enable the A and B flagellins or the sheath that contains them to be biosynthesized and, if this is the case, does not enable the *H. pylori* anchoring protein or the hook to be synthesized; and
- b) detecting an immunological reaction between the bacterial extract and antibodies directed against *H. pylori* and which are present in the sample.
- 50. The method as claimed in claim 49, wherein the aflagellate *H. pylori* strain also does not express the hook protein of *H. pylori*.
- 51. The method as claimed in claim 49, wherein the *H. pylori* bacterial strain is obtained from strain N6 having deposit Accession No. NCIMB 40512.
- 52. The method as claimed in claim 49, wherein the aflagellate strain of *H. pylori* is strain N6flbA having deposit Accession No. NCIMB 40747.
- 53. The method as claimed in claim 49, wherein the biological fluid is human serum, saliva, or urine.
- 54. The method as claimed in claim 49, wherein the immunological reaction is detected by Western blot or ELISA.
- 55. The method as claimed in claim 49, wherein the bacterial extract is a total bacterial extract of an aflagellate strain of H. pylori.

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- 56. The method as claimed in claim 49, wherein the bacterial extract is a n-octyl glucoside extract of strain N6flpA⁻ having deposit Accession No. NCIMB 40747.
- 57. The method as claimed in claim 43 or claim 49, wherein the *H. pylori* bacterial strain, or the bacterial extract of the said bacterial strain, is selected from the group consisting of:
 - (a) a bacterial strain lacking the hook protein of H. pylori;
 - (b) a recombinant bacterial strain obtained from the strain N6 (NCIMB 40512);
 - (c) a recombinant pacterial strain obtained from the strain N6 (NCIMB 40512) and lacking the hook protein of *H. pylori*;
 - (d) a recombinant bacterial strain obtained from the strain N6flbA⁻ (NCIMB 40747); or
 - (e) a recombinant bacterial strain obtained from the strain N6flbA⁻ (NCIMB 40747) and lacking the hook protein of *H. pylori*.
- 58. The method according to faim 57, wherein the bacterial extract is obtained after extracting with n-octyl glucoside.
- 59. The method according to claim 57, wherein the bacterial extract is obtained after extracting with PBS or with glycine.
 - 60. The method according to caim 57, wherein the flbA gene is SEQ ID NO:6.
- 61. The method according to claim 43 or claim 49, wherein the *flbA* gene is SEQ ID NO:6.--

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